

The mutant

Project N. \_\_\_\_\_  
 Block No. \_\_\_\_\_

Tag N. \_\_\_\_\_

Control → No plaques.  
 2899 → No plaques in either 10μl or 90μl  
 2904 → 10 plaques on 10μl  
 ~ 50 plaques on 90μl.

Infect 6 plaques from 2904 (F→Y) with 50μl DITSAF I &  
 in 2ml EG. → 6 hrs at 37°C.

Save cultures and plaques.

For 2899 : Re transform DITSAF I & without any dilution  
 — similarly transform with control

Overlay with top agar using 10μl & 90μl.  
 37°C O/N.

Control → 4 plaques (90μl)  
 2899 → 1 plaque (10μl)  
 13 plaques (90μl)

Save the plates at 4°C O/N.

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Issued & Understood by me,

*Handwritten signature*

Date

4/6/95

Invented by

Recorded by

*Handwritten signature*

Date

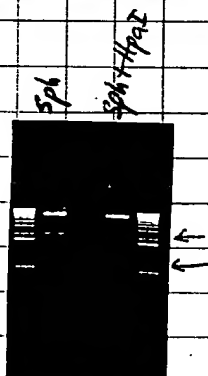
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Test F $\rightarrow$ Y mutants (6 clones)Standard mini prep.  
Digest with Hpa I.DNA was taken in 100  $\mu$ l TE.Hpa I  
digest

#1 seems to be correct clone.

Test #1 with sph and sph+Hp

If correct, sphI fragment with spl  
by about 600 bp.This gel proves that sphI fragment  
indeed shorter by about 600 bp.  $\frac{P}{2}$  $\therefore$  Thus #1 clone is correct and  
saved.

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